IN THE CLAIMS

This listing of claims provided below will replace all prior versions and listings of claims in the application.

Claims 1-33 (canceled).

- 34. (previously presented) A composition comprising florfenicol butyrate and a pharmaceutically acceptable solvent selected from N-methyl pyrrolidone, pyrrolidone, polyethylene glycol, propylene glycol, glycerol formal, isosorbid dimethyl ether, ethanol, dimethyl sulfoxide, tetrahydrofurfuryl alcohol, triacetin, and combinations thereof, wherein the composition is formulated as a composition for administration to a mammal by injection.
- 35. (previously presented) The composition of claim 34, wherein the solvent is selected from propylene glycol, glycerol formal, and combinations thereof.
- 36. (currently amended) The composition of claim 35, wherein the pharmaceutically acceptable <u>solvent</u> earrier is a combination of propylene glycol and glycerol formal.
- 37. (previously presented) The composition of claim 34, wherein the concentration of florfenical butvrate in the composition is at least about 200 mg/mL.

Claims 38-44 (canceled).

- 45. (previously presented) A method of treating a bacterial infection in a feline comprising injecting the feline with the pharmaceutical composition of claim 34.
- 46. (previously presented) A method of treating a bacterial infection in a feline comprising injecting the feline with the pharmaceutical composition of claim 35.
- 47. (previously presented) A method of treating a bacterial infection in a feline comprising injecting the feline with the pharmaceutical composition of claim 36.
- 48. (previously presented) A method of treating a bacterial infection in a feline comprising injecting the feline with the pharmaceutical composition of claim 37.

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- 49. (canceled).
- 50. (New) The pharmaceutical composition of claim 34, wherein the solvent is selected from the group consisting of propylene glycol, glycerol formal, N-methyl pyrrolidone, and combinations thereof.
- 51 (New) The pharmaceutical composition of claim 50, wherein the solvent is a combination of propylene glycol, glycerol formal, and N-methyl pyrrolidone.
- 52. (New) The pharmaceutical composition of claim 50, wherein the solvent is a combination of propylene glycol and N-methyl pyrrolidone.
- 53. (New) The pharmaceutical composition of claim 50, wherein the solvent is a combination of glycerol formal and N-methyl pyrrolidone.
- 54. (New) The pharmaceutical composition of claim 34, wherein the solvent is selected from the group consisting of tetrahydrofurfuryl alcohol, N-methyl pyrrolidone, and combinations thereof.
- 55. (New) The pharmaceutical composition of claim 54, wherein the solvent is a combination of tetrahydrofurfuryl alcohol and N-methyl pyrrolidone.
- 56. (New) The pharmaceutical composition of claim 34, wherein the solvent is selected from the group consisting of propylene glycol, glycerol formal, N-methyl pyrrolidone, polyethylene glycol, and combinations thereof.
- 57. (New) The pharmaceutical composition of claim 56, wherein the solvent is a combination of polyethylene glycol and N-methyl pyrrolidone.
- 58. (New) The pharmaceutical composition of claim 56, wherein the solvent is a combination of polyethylene glycol and glycerol formal.
- 59. (New) The pharmaceutical composition of claim 56, wherein the solvent is a combination of polyethylene glycol and propylene glycol.

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60. (New) The pharmaceutical composition of claim 56, wherein the solvent is a combination of polyethylene glycol, propylene glycol, and N-methyl pyrrolidone.

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